Characterization of the Human Gene for a Newly Discovered Carbonic Anhydrase, CA VII, and Its Localization to Chromosome 16

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Six carbonic anhydrase (CA) isozymes (CA I–VI) in mammals and other amniotes have been described. We have isolated an additional CA gene from a human genomic library and designated its putative product carbonic anhydrase VII (CA VII). The gene is approximately 10 kb long and contains seven exons and six introns found at positions identical to those determined for the previously described CA I, CA II, and CA III genes. The finding of a 17-bp GT-rich segment in a position 28 bp downstream of the poly(A)+ signal and the high correspondence of the 5' and 3' splice sites of the six introns with consensus junction sequences are consistent with the gene being functional. The 5' flanking regions of the CA VII gene do not contain the TATA and CAAT promoter elements usually found within 100 bp upstream of transcription initiation, but do contain a TTTAA sequence 102 nucleotides upstream of the initiation codon. The 5' region of the gene (−243 to +551) is GC-rich and contains 80 CpG dinucleotides and four possible Spl (GGGCGG or CCGCCC) binding sites. Northern analysis has identified the salivary gland as a major site of expression. The derived amino acid sequence of the CA VII gene is 263 amino acids long and has 50, 56, and 49% identity with human CA I, CA II, and CA III, respectively. No differences were found at any of the 39 positions that have remained invariant in all mammalian CA isozymes sequenced to date. Based on analysis of interspecific somatic cell hybrids, the human CA VII gene, CA7, was assigned to chromosome 16, with localization to the long arm at the q21–23 region by in situ hybridization. This is in contrast to the location of the CA I, CA II, and CA III gene cluster on human chromosome 8 and that of the human CA VI gene on chromosome 1.

INTRODUCTION

The carbonic anhydrases (CA; EC 4.2.1.1) are unusually diverse in their distribution for a multilocus enzyme family (Tashian, 1989). In mammals, six CA isozymes (CA I–CA VI) and two other partially characterized CA or CA-like isozymes, CA Y (= CA V?) and a CA-related protein (CARP), have been described (cf. Amor-Gueret and Levi-Strauss, 1990; Kato, 1990; Hewett-Emmett and Tashian, 1991). The expression of the CA genes ranges from their presence in certain cells of virtually all tissues (e.g., CA II) to their seemingly limited presence in one cell type of a specific tissue, e.g., CA VI in the serous accinar cells of the parotid and submandibular glands (Aldred et al., 1991) and CARP in the Purkinje cells of the cerebellum (Kato, 1990). Further diversity may be seen in their intracellular distribution where these CA isozymes may be cytosolic (CA I, CA II, CA III), membrane-bound (CA IV), mitochondrial (CA V), or present in secretory granules (CA VI) (cf. Tashian, 1989).

The CA isozyme genes vary in length from about 10 kb in human CA III to about 50 kb in human CA I, and they all contain seven coding exons that are interrupted by six introns at identical positions (cf. Hewett-Emmett and Tashian, 1991). The CA I gene, however, is unique in that it not only has an additional, noncoding exon, upstream of exon 1 (Fraser et al., 1989; Lowe et al., 1990), but also is under the control of two promoters that are apparently responsible for the expression of CA I either in red cells or in tissues other than red cells (Fraser et al., 1989; Tash-
ian et al., 1990). The CA I, CA II, and CA III genes are closely linked on chromosome 3 in the mouse (Beechey et al., 1990) and on the long arm of chromosome 8 in humans (Davis et al., 1987; Nakai et al., 1987). The human CA VI gene is located on the tip of the short arm on chromosome 1 (Sutherland et al., 1989).

During the course of screening a human genomic library with a mouse CA II cDNA probe, we isolated a CA-like gene (originally designated by us as CA Z), which on preliminary analysis appeared to code for a new CA isozyme. Here, we report the structure and chromosomal location of this new CA gene, the product of which we designate carbonic anhydrase VII (CA VII).

MATERIALS AND METHODS

Restriction Mapping

Physical maps of inserts from clones H10 and H11 were determined by single, double, and triple restriction digest analyses. Positioning of ambiguous sites was facilitated by hybridizing radiolabeled EcoRI fragments from the inserts to Southern blots of digested insert DNA. Restriction fragments from clones H21, 9-1B, and 2-4 were ordered and aligned by comparison to maps from clones H10 and H11. Certain subregions of some of the clones were mapped in greater detail by subcloning the fragment of interest prior to analysis of multiple digests with several additional enzymes.

DNA Blots

Restriction digests of genomic or cloned DNA were transferred from agarose gels to GeneScreen Plus (DuPont/NEN) by the alkaline method (Chomczynski and Qasba, 1984). Prehybridizations were carried out for 1 to 18 h at 42°C in 10–45% formamide, 5× SSPE (1× SSPE is 0.18 M NaCl, 10 mM NaH₂PO₄ (pH 7.4), 1.0 mM EDTA), 1.0% SDS, and 10% dextran sulfate. DNA probes were labeled by the mixed oligonucleotide random-priming technique (Hodgson and Fisk, 1987) and RNA probes were generated by incorporation of [α-³²P]UTP during SP6 transcription from linearized subclones in pSP64 or pSP65 vectors (Promega). Probes were added directly to the prehybridization solution for 18 to 48 h and washed initially at room temperature in two changes of 2× SSC, 0.1% SDS (1× SSC is 0.15 M NaCl, 0.015 M sodium citrate). Further washes with the same solution were performed at 55°C for low-stringency hybridization experiments, while higher-stringency hybridization filters were washed with at least two additional changes of 0.1× SSC, 0.1% SDS at 55–65°C. The filters were exposed to Kodak XAR-5 film at −70°C with intensifying screens. Blots were stripped for rehybridization by washing in 0.4 M NaOH and neutralizing with 0.2 M Tris–Cl (pH 7.5), 0.1× SSC, 0.1% SDS each for 1 h at 53°C.

Isolation of Genomic Clones

A chromosomal walk to isolate upstream clones overlapping the CA VII gene locus was carried out by screening 1× 10⁶ λ EMBL-3 recombinants (prepared by T. M. Johnson, this laboratory) made with partially digested DNA from an individual deficient in CA II (Sly et al., 1985). Standard plaque lifts (Maniatis et al., 1982) were hybridized to a probe derived from the most 5' EcoRI fragment of clones H10 and H11. Positive plaques were purified by two additional rounds of replating and screening. Phage DNA was prepared by polyethylene glycol precipitation of bacterial phage particles from 1 liter of liquid lysates generated by infection of Escherichia coli strain LE392 with 5 to 50× 10⁶ phage (Maniatis et al., 1982).

Subcloning and Sequencing

Fragments of interest were routinely subcloned into the pUC series of plasmid vectors [pUC8, pUC9 (Viera and Messing, 1982; Hanna et al., 1984); pUC18, pUC19 (Norrander et al., 1983)] or pBSM13+ or pBSM13− (Stratagene) using standard methods. A series of nested deletion subclones was created to facilitate sequencing the regions containing exons 5, 6, and 7 (Pontz et al., 1982; Henikoff, 1984). Fragments to be sequenced by the chemical cleavage method were end-labeled with T4 polynucleotide kinase and [γ-³²P]ATP, digested to create fragments with only one labeled end, and separated on polyacrylamide gels prior to base modification and cleavage reactions (Maxam and Gilbert, 1980). Double-stranded, supercoiled plasmids, prepared by the boiling method (Maniatis et al., 1982), were utilized for dideoxy chain termination sequencing (Chen and Seeburg, 1985; Hattori and Sakaki, 1986) with [α-³⁵S]dATP (Biggin et al., 1983) and the Sequenase T7 DNA polymerase (Tabor and Richardson, 1987). Ladders were resolved on 6% polyacrylamide/7.6 M urea buffer gradient gels (Biggin et al., 1983), which were fixed with 10% acetic acid and dried onto a glass plate that had been pretreated with γ-(methacryloxy)propyltrimethoxysilane (Garoff and Ansorge, 1981). Attempts were made to obtain sequence data from both strands, particularly within the exons. When this was not achieved, the same strand was sequenced at least twice. Sequences were compiled and analyzed with the aid of
the Fustell sequence analysis software (Fustell and Kafatos, 1982a,b, 1984).

**Chromosomal Mapping**

The 40 human–mouse somatic cell hybrid lines, which were derived from fusions of 14 unrelated human cell lines and 4 mouse cell lines, came from previously described hybrid sets (Shows et al., 1982, 1984; Sakaguchi and Shows, 1982). Each hybrid was analyzed for the presence or absence of the human CA VII fragment, as well as for the presence of individual human chromosomes, mapped enzyme markers, and in some cases mapped DNA probes (Shows et al., 1982, 1984, 1978; Shows, 1983). Southern blots of XbaI-digested DNA of the hybrids were hybridized to the 32P-radiolabeled 3.0-kb EcoRI fragment from clone H11 as described (Shows, 1983; Shows et al., 1984). Filters were washed twice for 30 min at 65°C with 0.1× SSC, 0.1% SDS and exposed to film for 1–3 days at −70°C with intensifying screens. Chromosomes for in situ hybridization were prepared from human lymphocytes by previously detailed methods (Dutrillaux and Viegas-Pequignot, 1981; Zabel et al., 1983) for obtaining long chromosomes. Hybridization to the same probe used above (3H-labeled), autoradiography, and G-banding were also as described (Zabel et al., 1983).

**RESULTS**

**Isolation and Identification of the Carbonic Anhydrase VII Gene**

The screening of the bacteriophage λ library has been described previously (Venta et al., 1984). Briefly, the insert from mouse CA II cDNA clone pMCAII (Curtis, 1983) was hybridized under reduced stringency conditions (35% formamide) to 3 × 108 recombinant phage. Six recombinants were plaque-purified from a total of 25 potential positive colonies. Clones H10 and H11, isolated from adjacent, weakly hybridizing plaques, had inserts with restriction maps unlike those of the other clones which were subsequently shown to contain the CA I or CA II genes (Bartworth et al., 1991; Venta et al., 1985). Although the inserts of these clones differ in length (15.1 kb for H10 vs 12.1 kb for H11), the configuration of restriction sites at the ends of the inserts is identical in both clones, suggesting that the H11 insert differs from H10 only by a 3-kb internal deletion (Fig. 1). The segment of DNA that cross-hybridized to the pMCAII probe was mapped to the center of the H10 insert within 5.1-kb BamHI, 1.3-kb EcoRI, and 6.9-kb HindIII fragments (Fig. 1). Fragments of identical size were detected in a Southern blot of human genomic DNA that was probed with a subfragment of the 7.3 kb EcoRI fragment from H10 (Fig. 2). This demonstrates that no major rearrangements had taken place during cloning within a minimum region of 12.0 kb in H10 that extends from the 5′ end of the 7.1-kb BamHI fragment to the 3′ end of the 6.9-kb HindIII fragment, and that H10, not H11, faithfully represents the arrangement of DNA in this region of the genome. The simplicity of the banding pattern indicates that a gene located here is probably present as a single copy in the haploid genome.

Sequence analysis of this region revealed an open reading frame, bounded by consensus acceptor and donor splice sites, that predicted an amino acid sequence which was 56, 65, and 52% identical to the corresponding regions of the human CA I, CA II, and CA III proteins, respectively. This region of the carbonic anhydrases is precisely encoded by exon 6 of all other characterized vertebrate CA genes (Tashian et al., 1990; Hewett-Emmett and Tashian, 1991). Additional potential coding material, which corresponded to a CA-like exon 3, was identified by reduced stringency hybridization (10% formamide) of H11 subfragments to a single-stranded RNA probe derived from the five exons of the mouse CA II cDNA (data not shown). While no additional independent regions of cross-hybridization were detected with this probe or with a probe derived from exon 7 and the 3′ untranslated region (UTR) of the mouse CA II gene, genomic or cDNA probes containing the first two exons of the human CA II gene did detect, under similar low-stringency conditions, a CA-like exon 2 in the 5.9-kb EcoRI and 7.2-kb HindIII fragments of 9-1B and 2-4, two clones obtained by genomic walking from H10 (data not shown). The inability of the mouse probe to hybridize to this region, which is present in H10 and H11, is probably a result of its slightly lower level of sequence identity with the identified exon compared to that of the human CA II gene (56% vs 60%) and a lower number of hydrogen bonds per base pair in the region of hybridization. The locations of exons 4, 5, and 7 were easily determined by directly sequencing the DNA downstream of exons 3 and 6, and a cluster of seven GC-dinucleotide-containing restriction sites (one NarI, three SmaI, one BssHII, and two SacII; data not shown) within a 0.5-kb region of an additional overlapping clone, H21, led to the discovery of exon 1 within a suspected “HTF island” (Lindsay and Bird, 1987), which is often a landmark for the 5′ end of transcribed genes (Bird, 1987).

**Nucleotide Sequence of the CA VII Gene**

The nucleotide sequence encompassing the CA VII gene region from 510 bp upstream of the putative
translation initiation codon in exon 1 to approximately 1.3 kb downstream of a single poly(A)\(^+\) addition signal is shown in Fig. 3. The seven exons and six introns define a gene of approximately 10 kb, assuming a 5' untranslated region (UTR) of about 70 nucleotides (see below) and that polyadenylation occurs near the addition signal (nucleotide 7152). The presence of a 17-nucleotide GT-rich segment (7185–7201) at a position (28 nucleotides downstream of the poly(A)\(^+\) signal) where similar sequences have been found to be necessary for the formation of proper 3' termini in various mRNAs (McLauchlan et al., 1985; Birnsteil et al., 1985) is consistent with the signal being functional. This suggests that the putative CA VII mRNA would be 1.5–1.6 kb long and would contain about 650 untranslated nucleotides at the 3' end.

The first six exons are interrupted by intervening sequences at positions that are identical to those seen in the CA I, CA II, and CA III genes (cf. Hewett-Emmett and Tashian, 1991). These introns contain invariant GT and AG dinucleotides at their 5' and 3' ends (Breathnach and Chambon, 1981) that are part of larger consensus donor and acceptor splice junctions (Mount, 1982). The high degree of sequence identity between these splice junctions and the consensus sequences shown in Table 1 is consistent with their being functional. A pentameric consensus sequence, CT(CG)A(GTC), which has been identified as the point at which the 5' end of many spliced introns forms a lariat branch (Ruskin et al., 1984), was found at the 3' end of introns 1, 2, 4, 5, and 6 within 21–29 nucleotides of the splice junction. Computer searches of a number of introns have led to the discovery of similar sequences that have been postulated to bind via hydrogen bonding with a conserved region of the U2 snRNA during the splicing process (Keller and Noon, 1984).
Four Alu repeat elements are located in the CA VII gene region (arrows in Fig. 1; boxed sequences in Fig. 3). One element within intron 5 (nucleotides 5208–5687) has been sequenced in its entirety and found to be flanked by identical 12-bp direct repeats. Three other Alu repeats have been partially sequenced. Two of these were found in intron 2 and represent the 3’ end of a repeat oriented in the same direction as the CA VII gene (nucleotides 3638–3689) and the middle of a repeat oriented in the opposite direction (nucleotides 3439–3637). Another oppositely oriented element was found in the 3’ UTR (nucleotides 7911–8204). A poly(GT) sequence, (GT)14, was found in the first intron (nucleotides 2151–2178).

The region of the CA VII gene upstream of the putative translation initiation codon does not contain a consensus TATA RNA polymerase II promoter element or a CAAT box element, both of which are commonly found approximately 20–30 and 40–100 bp, respectively, upstream of the start site of transcription for many eukaryotic genes (Breathnach and Chambon, 1981). Located at the −102 position relative to the initiation codon, however, is the sequence TTAAAA, which closely resembles the TTAA sequence found at the −27 position relative to the transcriptional start site of the human adenosine deaminase gene (Valercio et al., 1985), and the TTAAAA found at the −23 position relative to the start site of the human thrombospondin gene (Laherty et al., 1989). The former sequence is located in a GC-rich region upstream of the structural gene. The GC content of the 5’ end of the CA VII gene is also high. Calculated over ranges of 100 bases centered around each nucleotide from −243 to +551 relative to the putative start codon, GC content ranges from 65 to 84%. Eighty CpG dinucleotides, or an average of about 1 in 10 bp, are found in this region. Not unexpectedly, several GC boxes or potential Sp1 transcription factor binding sites (Kadonaga et al., 1986) were discovered in this region, including strict consensus sequences (GGGCGG or CCGCCC) at positions −185, −143, +155, and +398 relative to the start codon.

**The Predicted CA VII Protein.**

The protein predicted by the seven exons of the CA VII gene is 263 amino acids long. An alignment with the cytosolic human CA isozymes (CA I, II, and III) in Fig. 4 shows an extra amino acid at the N-terminus relative to the CA I protein and two extra relative to CA II and CA III: at the C-terminus, CA VII has two extra amino acids compared to CA I and one more than CA II and CA III. The only other gap needed to align the CA VII protein is found at residue 126 of CA II and CA III, where an extra amino acid is found in CA I and CA VII. In contrast, to maximally align the CA VII sequence with the predicted protein sequence of the human CA VI secreted salivary isozyme, a total of 11 gaps need to be introduced into the two sequences.

Overall, the predicted CA VII protein shares 50, 56, 49, and 37% sequence identity with human CA I, CA II, CA III, and CA VI, respectively. Sequence identity is not evenly distributed throughout the length of the protein, however. The alignment in Fig. 4 shows that there are distinct clusters of conservation and divergence between the CA VII protein and the other CAs. For example, residues 92–98, 104–111, 117–124, and 191–203 are strongly conserved regions among the CA I, CA II, CA III, and CA VI isozymes, and also remained unchanged for the most part in CA VII. In contrast, amino acids 150–170 are poorly conserved. The conserved regions of CA VII tend to fall into one or both of two categories. These residues usually either occupy positions that correspond to active site residues in human CA I and CA II (asterisks in Fig. 4) or are in positions where the residues are conserved in all mammalian carbonic anhydrase isozymes that have been sequenced to date (underlined in Fig. 4). Of 36 active site residues of CA VII, 33 are conserved in at least one other isozyme, including the key residues involved in the zinc ion ligands, His-94, His-96, and His-119, as well as Thr-199, the residue hydrogen-bonded to the zinc-bound solvent molecule (cf. Linds-kog, 1982; Silverman and Vincent, 1983). Only the active site residues at positions 69, 91, and 204 of CA VII are not found in the corresponding position in at least one of the other human isozymes. In each of these cases, however, the position is poorly conserved, as each human isozyme has its own unique amino acid. The active site changes in CA VII do represent conservative replacements with respect to one of the other isozymes: Asp for Glu at codon 69 of CA II and Lys for Arg at codon 91 of CA III. While a Ser at active site residue 204 is not found in any of the other human CAs, its presence in turtle CA I (Hewett-Emmett and Tashjian, 1991) suggests that this residue is not incompatible with enzyme function. There are no differences in the predicted protein sequence of CA VII at any of the 39 positions that are conserved in all vertebrate CA isozymes for which sequence information is currently available.

**Tissue-Specific Expression of the CA VII Gene.**

Figure 5A is an autoradiogram of a Northern blot of total RNA from various baboon tissues and several mammalian cell lines that was probed with a frag-
ment containing exon 3 from the CA VII gene. A single mRNA of approximately 1.6 kb migrates between the 18 S and 16 S markers in the salivary gland RNA (lane 8). When the same blot was stripped and reprobed with a human CA II cDNA insert (Fig. 5R), a similar sized mRNA was detected in the expected tissues: brain, kidney, liver, salivary gland, testes, and liver poly(A)⁺ RNA (lanes 2, 4, 5, 8, 10, and 16). The absence of a CA II mRNA in spleen (lane 9), lung (lane 6), and thyroid (lane 11) was unexpected but may be accounted for, at least in the latter two cases, by reduced levels of RNA on the membrane, since the background hybridization found in all the other lanes is absent in lanes 6 and 11 of Fig. 5A. The nature of the high-molecular-weight RNA hybridizing to the CA VII probe at the top of the HT1080 fibrosarcoma cell line lane (lane 13) is not known.

Chromosomal Location of the CA VII Gene

The 3.0-kb 5' EcoRI junction fragment from clone H11 was used as a hybridization probe to detect the presence of the human CA VII gene in human–mouse somatic cell hybrids. Figure 6 shows a representative Southern blot of XbaI-digested DNA from several hybrid cell lines. The probe detects a 12.0-kb fragment in human genomic DNA and an 8.6-kb fragment in mouse genomic DNA. Cross-hybridization of the mouse fragment was not due to the presence of exon 2 in the probe, but rather to a region that was found to reside in a 0.6-kb BglII/BamHI subfragment within the first intron (underlined sequence between nucleotides 1548 and 2109 in Fig. 3). In addition to human and mouse, this subfragment detects a single band in genomic DNA from macaque and rabbit (data not shown). The nature of this conserved region and its relationship to the CA VII gene are unclear. Computer analysis failed to show significant sequence similarity between the translated sequence of this fragment and the other carbonic anhydrase isozymes and search of the GenBank and EMBL databases revealed no identity with known genes. The presence of a pair of consensus acceptor and donor splice junction sequences, which would suggest an additional conserved exon in this region, was also not detected.

DNA from the somatic cell hybrids either had the mouse fragment only, and therefore lacked the human chromosome containing the human CA VII gene (Fig. 6, lanes 1–3), or had both human and mouse bands and therefore carried the human chromosome containing the CA VII gene (Fig. 6, lanes 4–6). The segregation analysis of the CA VII marker and various human chromosomes found in 40 somatic cell hybrids is shown in Table 2 and establishes the location of the CA VII gene on human chromosome 16. For each of the other human chromosomes, at least 21% of the hybrids were discordant.

In situ hybridization of the same probe to metaphase spreads of human lymphocyte chromosomes was performed to confirm the above result and to regionally assign the position of the CA VII gene. Based on the distribution of 331 grains on 200 metaphases examined, the CA VII gene was localized to the long arm of chromosomes 16 at the q21–23 region (Fig. 7B). Figure 7A shows that only this region has a signal above the background. The percentage of metaphases with a grain at 16q21–23 was 14%, while 14.2% of the total grains were on chromosome 16, with 68.1% of these grains located at 16q21–23 (Fig. 7A).

DISCUSSION

The overall structural organization of the human CA VII gene, consisting of seven exons with six introns interrupting putative coding sequences at the same nucleotides that mark exon boundaries in other CA genes, is similar to that seen for mammalian CA II and CA III genes. The identification of a 1.6-kb transcript is consistent with the predicted gene size of 10 kb, which is based on two assumptions. The first is the inferred position of the 3' terminus downstream of the poly(A)⁺ addition signal, which defines a rather large 3' untranslated region of greater than 600 nucleotides, similar to sizes seen in CA II genes. The second is the likely location of the transcriptional initiation site in the vicinity of the TTTAA pentamer,

FIG. 3. Nucleotide sequence of the CA VII gene. The sequence begins just downstream of the HindIII site with the 0.9-kb EcoRI fragment of clones 2-4 and 9-1B (c.f. Fig. 1). The nucleotides are numbered from the first base sequenced. The numbering does not take into account gaps in the sequence in intron 1, intron 2 (two gaps), and the 3' flanking region of the gene. Each exon is designated by encircled numbers at the beginning of the exon and the translated protein sequence appears below the nucleotide sequence. Amino acids are numbered below the protein sequence according to the CA I numbering system (cf. Ref. (25)). The putative ATG start and TGA stop codons are boxed at nucleotides 511 and 6533, respectively. A poly(A)⁺ addition signal is also boxed at nucleotide 7152. A GT-rich segment, similar to those found to be necessary for 3' end formation of other mRNAs, is underlined 26 bp downstream of the poly(A)⁺ signal. A conserved region in intron 1 is underlined and described in the text. Potential lariat branch points are doubly underlined upstream of the beginning of exons 2, 3, 5, 6, and 7. Located between nucleotides 2151 and 2179 is a (GT)₄₄ repeat element that is underlined with dashes. Alu repeats are boxed in intron 2 (two repeats), intron 5, and the 3' flanking region. The repeat in intron 5 is flanked by 12-bp direct repeats (dashed boxes). Sequence data for nucleotides 783–786 could not be determined.
Nucleotide Sequences at Exon/Intron Splice Junctions*

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Consensus

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* Capital letters indicate nucleotide matches with the consensus sequence. Lowercase letters indicate mismatches. Invariant GT and AG dinucleotides at the beginning and end of each intron are underlined.

Consensus sequences are from Ref. (44).

located 102 bp upstream of the putative start codon in exon 1. This is the only sequence motif in a very GC-rich region that resembles a TATA promoter. If the cap site is located 20–30 bp downstream, an untranslated region of about 70–80 nucleotides, consistent with the sizes of the 5’ UTRs of CA II and CA III genes, would precede the start codon. Although the methylation status of the CA VII gene has not been assessed, a high concentration of CpG dinucleotides in the region of the first exon suggests the presence of an HTF island (Bird, 1987). Consistent with this is the presence of four GC-box hexamers which are potential binding sites for the general transcription factor Sp1, an activator of transcription for a number of viral and cellular promoters (Kadonaga et al., 1986; Jones et al., 1986; Briggs et al., 1986). While the relevance of the Sp1 factor to regulation of CA VII gene expression remains to be established, the presence of these potential binding sites may be further evidence that this region contains the 5’ end of the gene.

On one hand, the predicted sequence of the putative CA VII protein shows a relatively high degree of divergence relative to that of the other human CA isozymes, but on the other hand it demonstrates the existence of a selective pressure for conservation of regions expected to be critical for proper enzyme activity. The similar pattern of conservation and divergence in CA II correlates well with the idea that the tertiary structure of CA VII will be similar to that of the human CA II molecule. Figure 4 shows that the region encoded by the 3’ half of exon 6, amino acids 191–222, is relatively conserved among the human CA isozymes, whereas the part of the protein encoded by the 5’ half of exon 6, amino acids 171–190, is much less conserved. Examination of the three-dimensional structures of human CA I, CA II, and bovine CA III (Eriksson, 1988) reveals that the former region is buried in the center of the molecule and forms part of the active site cavity, where substitutions would tend to be tolerated less in terms of maintenance of enzyme function. The latter region, however, is found at the external surface of the molecule in an area that can probably diverge much more freely without affecting function. Similar observations hold true for other areas of divergence. Poorly conserved regions of CA VII in exon 2 (residues 32–60), exon 3 (residues 89–103 and 112–116), exon 5 (residues 149–169), and exon 7 (residues 232–248) are all predicted to be located toward the surface of the CA VII protein and away from the active site based on their locations in the other CAs. Interestingly, all key active site residues, including the zinc-ion liganding histidines (His-94, His-96, and His-119) and the residues (Ser-29, Gln-92, Glu-106, His-107, Glu-117, Tyr-194, Thr-119, Trp-209, and Asn-244) thought to be involved in a hydrogen-bonding network with the zinc-bound solvent molecule (cf. Lindskog, 1982; Silverman and Vincent, 1983), are strictly conserved in CA VII, as they are in all CA isozymes. It would thus appear that conservation within regions of the CA VII protein is nonrandom, such that residues that are critical for CA enzyme activity are maintained. This is a strong indication that the CA VII gene encodes a functional protein.

Because it was immediately obvious that the CA VII gene could not encode the well-characterized cytosolic CA I, CA II, or CA III proteins, it was initially thought that its product could be either one of the less well described CA isozymes or a completely novel form of the enzyme. The latter possibility appears now to be most likely. The recently published cDNA sequence (Aldred et al., 1991) of the human secreted
HUMAN CARBONIC ANHYDRASE VII GENE

HCA VII

Comparison of the derived CA VII protein sequence with human CA I, CA II, CA III, and CA VI (cf. Refs. (1, 26)). Residues are shown for CA I, II, III, and VI that differ from CA VII in the corresponding position. Amino acids are numbered according to the CA I numbering system (cf. Ref. (26)). Gaps are introduced into the sequences to maintain the alignment of the residues and are indicated by dashes. Positions corresponding to active site residues in human CA I and CA II and bovine CA III are indicated by asterisks (cf. Ref. (65)), and positions that are conserved in all mammalian CA isozymes sequenced to date are underlined in the CA VII sequence (Refs. (2, 26, 33)). A potential N-linked glycosylation site at residues 71-73 is overlined. Intron positions for CA VII are indicated by numbered arrowheads. The single-letter amino acid code is used. Human CA VI has a leader sequence of 18 residues that is not shown (Ref. (1)).

salivary isozyme, CA VI, has eliminated this isozyme as a candidate for the CA VII gene product. Similarly, a comparison of the CA VII sequence with the partial protein sequence for the human CA IV membrane-bound CA isozyme (Zhu and Sly, 1990) shows only 33% identity (Tashian, 1989). No sequence data exist for the human CA V mitochondrial isozyme, but a comparison with the partial guinea pig CA V sequence (Hewett-Emmett et al., 1986) shows that the N-terminus of CA V aligns with amino acid 22 of other CA isozymes, implying that cleavage of an upstream mitochondrial import signal has occurred. It is unlikely

FIG. 5. Northern blot analysis of baboon and human RNAs with a CA VII probe in A containing exon 3 plus 43 bp of intron 2 and 116 bp of intron 3. The probe in B was the insert from a human CA II cDNA clone containing the entire cDNA except for the last 172 nucleotides of the 3' untranslated region (Ref. (43)). After hybridization, the blot was washed several times under low-stringency conditions, with a final high-stringency wash of 0.1X SSC, 0.1% SDS at 55°C. The RNAs in each lane are: 1, adrenal; 2, brain; 3, heart, 4, kidney; 5, liver; 6, lung; 7, pancreas; 8, salivary gland; 9, spleen; 10, testis; 11, thyroid; 12, thymus; 13, human fibrosarcoma HT 1080 cell line; 14, rat hepatoma HPC PA1 cell line; 15, human hepatoma HEP-G2 tube (10 mg); 16, human liver poly(A)+ RNA (2 μg).

FIG. 6. Representative Southern blot of human-mouse somatic cell hybrid DNA. The 3.0-kb EcoRI fragment from clone H11 hybridized to a single unique fragment in XhoI-digested DNA from mouse (8.6 kb, lane 7) and human (12.0 kb, lane 8). Cell hybrids missing the human specific CA VII fragment are in lanes 1-3 and were scored negative for the presence of the human chromosome containing the CA VII gene. Hybrids in lanes 4-6 contain the human fragment and were scored as positives.
TABLE 2
Segregation of the Carbonic Anhydrase VII Gene with Human Chromosomes in Human–Mouse Hybrids

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<td>60</td>
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</table>

* Concordant hybrids have either retained (+/+ or lost (−/−) the human CA VII signal together with a specific human chromosome.

b Discordant hybrids either retained the gene, but not a specific chromosome (+/−), or the reverse (−/+).

A 0% discordancy is the basis for chromosomal assignment.

that the N-terminus of the CA VII protein acts as such a signal because the first 22 amino acids of the CA VII protein align well with those of other CAs, and because the absence of positive charges in the first 18 residues and the presence of negatively charged residues at positions 10 and 11 are features not generally found in known import signals (Douglas et al., 1986).

In addition, the recently reported cDNA sequences of two distinct CA or CA-like proteins from mouse liver (Amor-Gueret and Levi-Strauss, 1991) and cerebellum (Kato, 1991) showed only low sequence identities (i.e., 52 and 37%, respectively) to the human CA VII sequence. This degree of conservation is in contrast to the much higher percentage identities (70–80%) seen when the orthologous gene products for other carbonic anhydrases, such as CA I, CA II, and CA III, are compared between these two species (Hewett-Emmett and Tashian, 1991) and suggests, therefore, that CA VII is not the human counterpart to either of these two mouse proteins.

Although the CA VII protein, unlike the cytosolic CA I, CA II, and CA III isozymes, has a single potential N-linked glycosylation site at codons 71–73, Asn–Asp–Ser, the above analysis suggests that it may be nonfunctional. The similar sizes of the predicted CA VII protein and the cytosolic enzymes, the higher degree of sequence conservation between it and these isozymes compared to that between it and the secreted or membrane-bound forms, and the apparent lack of a signal peptide all point to a probable cytosolic localization for CA VII, in which case it would not be glycosylated.

The detection of a band of the predicted size for a CA VII transcript (1.6 kb) in salivary gland RNA provides clear evidence that this locus is expressed, thus confirming predictions that the gene encodes a functional molecule. These predictions were based on findings of a high degree of widespread divergence in CA VII, except for a remarkably strict conservation of residues necessary for enzyme function. This high degree of divergence, which prevents other CA gene sequences from readily cross-hybridizing to CA VII sequences and vice versa (see Fig. 2), coupled with the detection of a band in RNA from only a single tissue, eliminates the probability that the CA VII probe is detecting a transcript from one of the other isozymes. Carbonic anhydrases I, II, and III have been detected in subcellular locations within various cell types of human, rat, and mouse salivary glands (Hennigar et al., 1983; Ikejima and Ito, 1984; Carpentier et al., 1985; Noda et al., 1986; Parkkila et al., 1990; Spicer et al., 1990; Aldred et al., 1991; Fernley et al., 1991).

However, since activity assays only detect CA activity and immunological methods can be subject to cross-reactivity, the presence of novel CA isozymes can be missed. For example, it is now clear that CA activity found in secretory granules of these glands is actually the CA VI isozyme. The additional presence of the CA VII isozyme is thus possible. Further work will be necessary to confirm this and to elucidate the roles of possibly five forms of carbonic anhydrase (CA I, CA II, CA III, CA VI, and CA VII) in salivary glands.

The finding that the CA VII gene is located on chromosome 16 and a recent report localizing the CA VI gene to human chromosome 1 (Sutherland et al., 1989) are the first examples of CA isozyme genes not linked to the CA1/CA3/CA2 gene cluster on human chromosome 8 (Davis et al., 1987; Nakai et al., 1987). Phylogenetic analyses indicate that the duplications giving rise to the CA I, CA II, and CA III genes occurred after the duplication that formed the CA VII gene (Hewett-Emmett and Tashian, 1991), suggest-
FIG. 7. Mapping of the CA VII gene to human chromosome 16 by in situ hybridization. (A) The total grain distribution after hybridization of the 3.0-kb EcoRI fragment from clone H11 to normal human lymphocyte metaphase chromosomes. Accumulation above background occurs only on the long arm of chromosome 16. (B) An idiogram localizing the majority of grains to 16q21–23.

ing that the latter gene may have dispersed prior to the formation of the former genes. The location of the CA VII gene on the long arm of chromosome 16 may be useful in further defining the linkage map of that portion of the genome (Keith et al., 1987). The polymorphic nature of poly(GT) repeats (Weber and May, 1989) and the 3' ends of Alu repeat elements (Hobbs et al., 1985) may facilitate the identification of informative CA VII probes that can be used for this purpose. The locations of these repeated elements may also be useful in identifying the presence of the CA VII gene region in previously cloned segments of chromosome 16 (Hildebrand et al., 1988).

The haptoglobin, adenine phosphoribosyltransferase, mitochondrial glutamate oxaloacetate transaminase, metallothionein I and II, lecithin–cholesterol acyltransferase, uvomorulin, and tyrosine aminotransferase genes, all located on the long arm of human chromosome 16, form part of a syntenic group that maps to mouse chromosome 8 (Nadeau, 1989; Lalley et al., 1989). Because of its location among these genes on human chromosome 16, it is expected
that the CA VII gene will also be found on chromosome 8 of the mouse.

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